

Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

1. (Currently Amended) An assay for determining level of amount of prostacyclin in a plasma sample comprising:

(1) providing a plasma sample on a surface coated with an anti-immunoglobulin antibody;

(2) incubating the plasma sample with an effective amount of an anti-6-keto-prostaglandin F_{1α} (6-keto-PGF_{1α}) antibody[[;]] wherein the [[an]] anti-immunoglobulin antibody that binds to the anti-6-keto-PGF_{1α}-antibody; and a conjugate comprising 6-keto-PGF_{1α} covalently bound to an aequorin mutant;

wherein said aequorin mutant comprises serine substitutions substitutions for all three cysteine residues as present in wild-type aequorin (Cys → Ser), wherein said aequorin mutant further comprises a single cysteine residue substituted at amino acid position 69 (Ala69 → Cys), 70 (Gly70 → Cys), 74(Gly74 → Cys) or 76 (Glu76 → Cys), and

wherein the 6-keto-PGF_{1α} is coupled to the aequorin mutant via reaction with the sulphydryl group of the single cysteine;

(3) removing any unbound anti-6-keto-PGF_{1α}-antibody and said conjugate from the plasma sample following incubation; and

(4) measuring and correlating light intensity of the 6-keto-PGF_{1α} bound to the anti-immunoglobulin antibody plasma sample; and with amount of prostacyclin within the plasma sample

(5) correlating the light intensity of the bound 6-keto-PGF_{1α} plasma sample, with the amount of prostacyclin in the plasma sample.

2. (Cancelled)

3. (Cancelled).

4. (Original) The assay of claim 1 wherein the plasma sample is obtained from a patient receiving intravenous prostaglandin therapy.

5. (Currently Amended) The assay of claim 1 wherein ~~the concentration of said conjugate in the assay~~ has a concentration of [[is]] about 1×10^{-10} M.

6. (Cancelled).

7. (Cancelled).

8. (Currently Amended) A method of determining an appropriate dose of amount of prostaglandin in a plasma sample for the treatment of primary pulmonary hypertension in a patient comprising:

- (1) providing a plasma sample from the patient on a surface coated with an anti-immunoglobulin antibody;
- (2) incubating the plasma sample with an effective amount of an anti-6-keto-prostaglandin F_{1α} (6-keto-PGF_{1α}) antibody[[;]] wherein the [[an]] anti-immunoglobulin antibody that binds to the anti-6-keto-PGF_{1α}-antibody; and a conjugate comprising 6-keto-PGF_{1α} covalently bound to an aequorin mutant;

wherein said aequorin mutant comprises serine substitutions substitutions for all three cysteine residues as present in wild-type aequorin (Cys → Ser), wherein said aequorin mutant further comprises a single cysteine residue substituted at amino acid position 69 (Ala69 → Cys), 70 (Gly70 → Cys), 74(Gly74 → Cys) or 76 (Glu76 → Cys), and

wherein the 6-keto-PGF_{1α} is coupled to the aequorin mutant via reaction with the sulphydryl group of the single cysteine,

- (3) removing any unbound anti-6-keto-PGF_{1α}-antibody and said conjugate from the plasma sample following incubation; and

(4) measuring and correlating light intensity of the amount of detected 6-keto- PGF_{1α} bound to the anti-immunoglobulin antibody; and

(5) correlating the light intensity of the bound 6-keto- PGF_{1α} with the appropriate dosage of prostaglandin for the patient amount of prostaglandin in the plasma sample.

9. (Cancelled).

10. (Cancelled).

11. (Currently Amended) The assay method of claim 8 wherein the plasma sample is obtained from a patient receiving intravenous prostaglandin therapy.

12. (Currently Amended) The assay method of claim 8 wherein the concentration of said conjugate in the assay is about 1×10^{-10} M.

13-21. (Cancelled)

22. (Currently Amended) A kit for measuring prostacyclin in plasma comprising:

- (1) an anti-6-keto-prostaglandin F_{1α} (6-keto-PGF_{1α}) antibody;
- (2) an anti-immunoglobulin antibody that binds to the anti-6-keto-PGF_{1α} antibody; and
- (3) a conjugate comprising 6-keto-PGF_{1α} covalently bound to an aequorin mutant; wherein said aequorin mutant comprises serine substitutions substitutions for all three cysteine residues as present in wild-type aequorin (Cys → Ser), wherein said aequorin mutant further comprises a single cysteine residue substituted at amino acid position 69 (Ala69 → Cys), 70 (Gly70 → Cys), 74(Gly74 → Cys) or 76 (Glu76 → Cys), and wherein the 6-keto-PGF_{1α} is coupled to the aequorin mutant via reaction with the sulphhydryl group of the single cysteine.